

Christian Heinis

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/2386308/publications.pdf>

Version: 2024-02-01

87
papers

5,159
citations

109321

35
h-index

88630

70
g-index

93
all docs

93
docs citations

93
times ranked

5367
citing authors

#	ARTICLE	IF	CITATIONS
1	An Engineered Protein Tag for Multiprotein Labeling in Living Cells. <i>Chemistry and Biology</i> , 2008, 15, 128-136.	6.0	940
2	Phage-encoded combinatorial chemical libraries based on bicyclic peptides. <i>Nature Chemical Biology</i> , 2009, 5, 502-507.	8.0	595
3	Cyclic peptide therapeutics: past, present and future. <i>Current Opinion in Chemical Biology</i> , 2017, 38, 24-29.	6.1	518
4	Bicyclic Peptide Inhibitor Reveals Large Contact Interface with a Protease Target. <i>ACS Chemical Biology</i> , 2012, 7, 817-821.	3.4	156
5	Directed evolution of O6-alkylguanine-DNA alkyltransferase for applications in protein labeling. <i>Protein Engineering, Design and Selection</i> , 2006, 19, 309-316.	2.1	136
6	Strategies to prolong the plasma residence time of peptidedrugs. <i>MedChemComm</i> , 2010, 1, 319-324.	3.4	130
7	Phage Selection of Cyclic Peptides for Application in Research and Drug Development. <i>Accounts of Chemical Research</i> , 2017, 50, 1866-1874.	15.6	117
8	Cyclization of peptides with two chemical bridges affords large scaffold diversities. <i>Nature Chemistry</i> , 2018, 10, 715-723.	13.6	113
9	Tools and rules for macrocycles. <i>Nature Chemical Biology</i> , 2014, 10, 696-698.	8.0	105
10	Peptide Ligands Stabilized by Small Molecules. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 1602-1606.	13.8	103
11	Acylated heptapeptide binds albumin with high affinity and application as tag furnishes long-acting peptides. <i>Nature Communications</i> , 2017, 8, 16092.	12.8	101
12	Encoded libraries of chemically modified peptides. <i>Current Opinion in Chemical Biology</i> , 2015, 26, 89-98.	6.1	99
13	Structurally Diverse Cyclisation Linkers Impose Different Backbone Conformations in Bicyclic Peptides. <i>ChemBioChem</i> , 2012, 13, 1032-1038.	2.6	81
14	Bicyclic Peptide Ligands Pulled out of Cysteine-Rich Peptide Libraries. <i>Journal of the American Chemical Society</i> , 2013, 135, 6562-6569.	13.7	78
15	Boosting the Sensitivity of Ligand-Protein Screening by NMR of Long-Lived States. <i>Journal of the American Chemical Society</i> , 2012, 134, 11076-11079.	13.7	75
16	Dithiol amino acids can structurally shape and enhance the ligand-binding properties of polypeptides. <i>Nature Chemistry</i> , 2014, 6, 1009-1016.	13.6	73
17	Measuring In Vivo Protein Half-Life. <i>Chemistry and Biology</i> , 2011, 18, 805-815.	6.0	71
18	Engineering Substrate Specificity of O6-Alkylguanine-DNA Alkyltransferase for Specific Protein Labeling in Living Cells. <i>ChemBioChem</i> , 2005, 6, 1263-1269.	2.6	68

#	ARTICLE	IF	CITATIONS
19	Phage Selection of Photoswitchable Peptide Ligands. <i>Journal of the American Chemical Society</i> , 2014, 136, 5880-5883.	13.7	67
20	Bicyclic Peptides with Optimized Ring Size Inhibit Human Plasma Kallikrein and its Orthologues While Sparing Paralogous Proteases. <i>ChemMedChem</i> , 2012, 7, 1173-1176.	3.2	66
21	De novo development of proteolytically resistant therapeutic peptides for oral administration. <i>Nature Biomedical Engineering</i> , 2020, 4, 560-571.	22.5	65
22	Phage selection of bicyclic peptides. <i>Methods</i> , 2013, 60, 46-54.	3.8	64
23	A Synthetic Factor Xlla Inhibitor Blocks Selectively Intrinsic Coagulation Initiation. <i>ACS Chemical Biology</i> , 2015, 10, 1861-1870.	3.4	64
24	Polycyclic Peptide Therapeutics. <i>ChemMedChem</i> , 2013, 8, 377-384.	3.2	63
25	Phage Selection of Chemically Stabilized α -Helical Peptide Ligands. <i>ACS Chemical Biology</i> , 2016, 11, 1422-1427.	3.4	63
26	Cyclic peptide FXII inhibitor provides safe anticoagulation in a thrombosis model and in artificial lungs. <i>Nature Communications</i> , 2020, 11, 3890.	12.8	61
27	Improving Binding Affinity and Stability of Peptide Ligands by Substituting Glycines with α -Amino Acids. <i>ChemBioChem</i> , 2013, 14, 1316-1322.	2.6	56
28	Identification of target-binding peptide motifs by high-throughput sequencing of phage-selected peptides. <i>Nucleic Acids Research</i> , 2014, 42, e169-e169.	14.5	55
29	Development of a Selective Peptide Macrocyclic Inhibitor of Coagulation Factor XII toward the Generation of a Safe Antithrombotic Therapy. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 3742-3746.	6.4	53
30	Synthesis and Photochemical Properties of Oligo-ortho-azobenzenes. <i>Journal of Organic Chemistry</i> , 2011, 76, 9826-9834.	3.2	48
31	Bicyclization and Tethering to Albumin Yields Long-Acting Peptide Antagonists. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 10187-10197.	6.4	47
32	Cys-Cys and Cys-Lys Stapling of Unprotected Peptides Enabled by Hypervalent Iodine Reagents. <i>Angewandte Chemie - International Edition</i> , 2021, 60, 9022-9031.	13.8	47
33	Peptide Macrocyclic Inhibitor of Coagulation Factor XII with Subnanomolar Affinity and High Target Selectivity. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 1151-1158.	6.4	45
34	Phage display libraries of differently sized bicyclic peptides. <i>MedChemComm</i> , 2013, 4, 145-150.	3.4	42
35	Precisely Regulated and Efficient Locking of Linear Peptides into Stable Multicyclic Topologies through a One-Pot Reaction. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 4458-4463.	13.8	39
36	Development of Potent and Selective <i>S. aureus</i> Sortase A Inhibitors Based on Peptide Macrocycles. <i>ACS Medicinal Chemistry Letters</i> , 2016, 7, 606-611.	2.8	37

#	ARTICLE	IF	CITATIONS
37	Thiol-to-amine cyclization reaction enables screening of large libraries of macrocyclic compounds and the generation of sub-kilodalton ligands. <i>Science Advances</i> , 2019, 5, eaaw2851.	10.3	30
38	Enzymatic Cyclisation of Peptides with a Transglutaminase. <i>ChemBioChem</i> , 2011, 12, 38-42.	2.6	29
39	Post-translational modification of genetically encoded polypeptide libraries. <i>Current Opinion in Chemical Biology</i> , 2011, 15, 355-361.	6.1	28
40	Phage selection of cyclic peptide antagonists with increased stability toward intestinal proteases. <i>Protein Engineering, Design and Selection</i> , 2013, 26, 81-89.	2.1	28
41	Phage Selection of Peptide Macrocyces against β -Catenin To Interfere with Wnt Signaling. <i>ChemMedChem</i> , 2016, 11, 834-839.	3.2	28
42	Combination of polycarboxybetaine coating and factor XII inhibitor reduces clot formation while preserving normal tissue coagulation during extracorporeal life support. <i>Biomaterials</i> , 2021, 272, 120778.	11.4	28
43	Chemical Macrocyclization of Peptides Fused to Antibody Fc Fragments. <i>Bioconjugate Chemistry</i> , 2012, 23, 1856-1863.	3.6	27
44	Phage selection of bicyclic peptides binding Her2. <i>Tetrahedron</i> , 2014, 70, 7733-7739.	1.9	27
45	Evolving the Substrate Specificity of O ⁶ -Alkylguanine-DNA Alkyltransferase through Loop Insertion for Applications in Molecular Imaging. <i>ACS Chemical Biology</i> , 2006, 1, 575-584.	3.4	25
46	Phage Selection of Bicyclic Peptide Ligands of the Notch1 Receptor. <i>ChemMedChem</i> , 2015, 10, 1754-1761.	3.2	25
47	Bicyclic Peptides Conjugated to an Albumin-Binding Tag Diffuse Efficiently into Solid Tumors. <i>Molecular Cancer Therapeutics</i> , 2015, 14, 151-161.	4.1	25
48	Engineered Peptide Macrocyces Can Inhibit Matrix Metalloproteinases with High Selectivity. <i>Angewandte Chemie - International Edition</i> , 2019, 58, 11801-11805.	13.8	23
49	Polar Hinges as Functionalized Conformational Constraints in (Bi)cyclic Peptides. <i>ChemBioChem</i> , 2017, 18, 387-395.	2.6	18
50	Improving the Binding Affinity of in vitro Evolved Cyclic Peptides by Inserting Atoms into the Macrocycle Backbone. <i>ChemBioChem</i> , 2016, 17, 2299-2303.	2.6	17
51	Phage Selection of Bicyclic Peptides Based on Two Disulfide Bridges. <i>Methods in Molecular Biology</i> , 2015, 1248, 119-137.	0.9	17
52	Pattern-Based Sensing of Peptides and Aminoglycosides with a Single Molecular Probe. <i>Organic Letters</i> , 2013, 15, 3456-3459.	4.6	16
53	Precisely Regulated and Efficient Locking of Linear Peptides into Stable Multicyclic Topologies through a One-Pot Reaction. <i>Angewandte Chemie</i> , 2017, 129, 4529-4534.	2.0	15
54	Picomole-Scale Synthesis and Screening of Macrocyclic Compound Libraries by Acoustic Liquid Transfer. <i>Angewandte Chemie - International Edition</i> , 2021, 60, 21702-21707.	13.8	14

#	ARTICLE	IF	CITATIONS
55	Synthesis and direct assay of large macrocycle diversities by combinatorial late-stage modification at picomole scale. <i>Nature Communications</i> , 2022, 13, .	12.8	14
56	Generation of a Large Peptide Phage Display Library by Self-Ligation of Whole-Plasmid PCR Product. <i>ACS Chemical Biology</i> , 2020, 15, 2907-2915.	3.4	13
57	Cysâ€Cys and Cysâ€Lys Stapling of Unprotected Peptides Enabled by Hypervalent Iodine Reagents. <i>Angewandte Chemie</i> , 2021, 133, 9104-9113.	2.0	13
58	Synthesis of DNAâ€Encoded Disulfideâ€and Thioetherâ€Cyclized Peptides. <i>ChemBioChem</i> , 2020, 21, 543-549.	2.6	12
59	Macrocycle synthesis strategy based on step-wise â€adding and reactingâ€three components enables screening of large combinatorial libraries. <i>Chemical Science</i> , 2020, 11, 7858-7863.	7.4	12
60	Screening of Large Molecule Diversities by Phage Display. <i>Chimia</i> , 2011, 65, 843-845.	0.6	10
61	Directed Evolution of Bicyclic Peptides for Therapeutic Application. <i>Chimia</i> , 2013, 67, 910-915.	0.6	10
62	Tracking chemical reactions on the surface of filamentous phage using mass spectrometry. <i>Chemical Communications</i> , 2014, 50, 5267-5269.	4.1	9
63	Development of Selective FXIa Inhibitors Based on Cyclic Peptides and Their Application for Safe Anticoagulation. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 6802-6813.	6.4	8
64	Bicyclic Peptide Antagonists Derived from Genetically Encoded Combinatorial Libraries. <i>Chimia</i> , 2011, 65, 677-679.	0.6	7
65	Generation of a 100-billion cyclic peptide phage display library having a high skeletal diversity. <i>Protein Engineering, Design and Selection</i> , 2021, 34, .	2.1	7
66	Cyclative Release Strategy to Obtain Pure Cyclic Peptides Directly from the Solid Phase. <i>ACS Chemical Biology</i> , 2022, 17, 181-186.	3.4	7
67	Two General Methods for the Isolation of Enzyme Activities by Colony Filter Screening. <i>Chemistry and Biology</i> , 2002, 9, 383-390.	6.0	6
68	Using Peptide Loop Insertion Mutagenesis for the Evolution of Proteins. <i>Methods in Molecular Biology</i> , 2010, 634, 217-232.	0.9	5
69	Measuring net protease activities in biological samples using selective peptidic inhibitors. <i>Analytical Biochemistry</i> , 2012, 427, 18-20.	2.4	5
70	Tissue Factor-Independent Coagulation Correlates with Clinical Phenotype in Factor XI Deficiency and Replacement Therapy. <i>Thrombosis and Haemostasis</i> , 2021, 121, 150-163.	3.4	5
71	Solid-phase peptide synthesis on disulfide-linker resin followed by reductive release affords pure thiol-functionalized peptides. <i>Organic and Biomolecular Chemistry</i> , 2022, 20, 5699-5703.	2.8	5
72	Chemical Biology Approaches to Membrane Homeostasis and Function. <i>Chimia</i> , 2011, 65, 849-852.	0.6	3

#	ARTICLE	IF	CITATIONS
73	Engineered Peptide Macrocycles Can Inhibit Matrix Metalloproteinases with High Selectivity. <i>Angewandte Chemie</i> , 2019, 131, 11927-11931.	2.0	3
74	Towards the Development of Orally Available Peptide Therapeutics. <i>Chimia</i> , 2021, 75, 514.	0.6	3
75	Combining biological and chemical diversity. <i>Nature Chemistry</i> , 2021, 13, 512-513.	13.6	3
76	Bypassing bacterial infection in phage display by sequencing DNA released from phage particles. <i>Protein Engineering, Design and Selection</i> , 2017, 30, 761-768.	2.1	2
77	A releasable disulfide-linked peptide tag facilitates the synthesis and purification of short peptides. <i>Chemical Communications</i> , 2020, 56, 2917-2920.	4.1	2
78	In Vitro-Evolved Peptides Bind Monomeric Actin and Mimic Actin-Binding Protein Thymosin- β 4. <i>ACS Chemical Biology</i> , 2021, 16, 820-828.	3.4	2
79	Picomole-scale Synthesis and Screening of Macrocyclic Compound Libraries by Acoustic Liquid Transfer. <i>Angewandte Chemie</i> , 2021, 133, 21870-21875.	2.0	2
80	Chemical biology & drug discovery. <i>European Journal of Medicinal Chemistry</i> , 2014, 88, 1-2.	5.5	1
81	The Partnership of DMCCB and LS2. <i>Chimia</i> , 2018, 72, 817-818.	0.6	1
82	Phage Display Selected Cyclic Peptide Inhibitors of Kallikrein-Related Peptidases 5 and 7 and Their <i>In Vivo</i> Delivery to the Skin. <i>Journal of Medicinal Chemistry</i> , 2022, 65, 9735-9749.	6.4	1
83	Fast Directed Evolution of Non-Immunoglobulin Proteins by Somatic Hypermutation in Immune Cells. <i>ChemBioChem</i> , 2005, 6, 804-806.	2.6	0
84	The 4th Young Faculty Meeting – Science and Funding in its Different Varieties. <i>Chimia</i> , 2011, 65, 818-820.	0.6	0
85	Innenrücktitelbild: Precisely Regulated and Efficient Locking of Linear Peptides into Stable Multicyclic Topologies through a One-Pot Reaction (<i>Angew. Chem.</i> 16/2017). <i>Angewandte Chemie</i> , 2017, 129, 4701-4701.	2.0	0
86	Drugs Based on de novo-developed Peptides are Coming of Age <i>Chimia</i> , 2018, 72, 426-427.	0.6	0
87	Chemical Biology and Drug Discovery Symposium at the LS2 Annual Meeting 2021. <i>Chimia</i> , 2021, 75, 342-342.	0.6	0